FILE 'HOME' ENTERED AT 11:57:42 ON 29 NOV 2000

=> file caplus medline embase wpids biosis

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FULL ESTIMATED COST

0.15 0.15

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FILE 'WPIDS' ENTERED AT 11:58:21 ON 29 NOV 2000 COPYRIGHT (C) 2000 DERWENT INFORMATION LTD

FILE 'BIOSIS' ENTERED AT 11:58:21 ON 29 NOV 2000 COPYRIGHT (C) 2000 BIOSIS(R)

=> s phytostenol# or phytosterol# or sitostenol# or sitostenol# or sitosterol#

L1 18440 PHYTOSTENOL# OR PHYTOSTEROL# OR SITOSTENOL# OR SITOSTENOL# OR SITOSTEROL#

=> s fatty(w)acid#

L2 535195 FATTY(W) ACID#

=> s unsaturat?(w)fatty(w)acid#

L3 35497 UNSATURAT?(W) FATTY(W) ACID#

=> s 11 and 13

L4 225 L1 AND L3

=> s hypocholest? or lower(n)cholest? or reduct?(s)cholest?

L5 53398 HYPOCHOLEST? OR LOWER(N) CHOLEST? OR REDUCT?(S) CHOLEST?

=> s 14 and 15

L6 11 L4 AND L5

=> d kwic ibib so 16 1-11

L6 ANSWER 1 OF 11 CAPLUS COPYRIGHT 2000 ACS

TI Method for manufacturing fat-soluble **phytosterol** or phytostanol ester of **unsaturated fatty acid**

AB The present invention provides a method for manufg. fat-sol.

phytosterol or phytostanol ester of unsatd.

fatty acid for inhibiting the absorption of cholesterol

and foodstuffs contg. the same. The method for manufg. fat-sol.

```
phytosterol or phytostanol ester of unsatd.
     fatty acid comprises the steps of: esterification of
     phytosterol or phytostanol with unsatd. fatty
     acid by dissolving them in a nonpolar org. solvent with a basic
     catalyst and adding a carboxyl group activating agent dissolved.
ST
     phytosterol fatty ester fat soluble prodn cholesterol
     absorption redn; phytostanol fatty ester fat soluble prodn
     cholesterol absorption redn; sitosterol oleate
     prodn esterification cholesterol lowering effect
     Fatty acids, preparation
     RL: BUU (Biological use, unclassified); IMF (Industrial manufacture);
BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (esters; method for manufg. fat-sol. phytosterol or
        phytostanol ester of unsatd. fatty acid)
IT
     57-88-5, Cholesterol, biological studies
     RL: BPR (Biological process); BIOL (Biological study); PROC (Process)
        (method for manufg. fat-sol. phytosterol or phytostanol ester
        of unsatd. fatty acid)
ΙT
     3712-16-1P, .beta.-Sitosterol oleate
     RL: BUU (Biological use, unclassified); IMF (Industrial manufacture);
BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (method for manufg. fat-sol. phytosterol or phytostanol ester
        of unsatd. fatty acid)
ΤТ
     83-46-5, .beta.-Sitosterol
                                  112-80-1, Oleic acid, reactions
     RL: RCT (Reactant)
        (method for manufg. fat-sol. phytosterol or phytostanol ester
        of unsatd. fatty acid)
ACCESSION NUMBER:
                         2000:742194 CAPLUS
DOCUMENT NUMBER:
                         133:323278
TITLE:
                         Method for manufacturing fat-soluble
                       phytosterol or phytostanol ester of
                       unsaturated fatty acid
INVENTOR(S):
                         Chung, Dae-Won; Noh, Seung-Kwon; Kim, Kab-Sig
PATENT ASSIGNEE(S):
                         Eugene Science, Inc., S. Korea
SOURCE:
                         PCT Int. Appl., 19 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
     WO 2000061694
     PATENT NO. KIND DATE
                                           -----
                     A1 20001019 WO 1999-KR569 19990921
     WO 2000061694
         W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ,
             DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ,
             TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
             DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
             CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
PRIORITY APPLN. INFO.:
                                           KR 1999-12965
                                                             19990413
     PCT Int. Appl., 19 pp.
     CODEN: PIXXD2
REFERENCE COUNT:
REFERENCE(S):
                         (1) Henkel; WO 9925362 A1 1999 CAPLUS
                         (2) Raisio Benecol Oy; WO 9956558 Al 1999 CAPLUS
                         (3) Raision Margarini; WO 9219640 Al 1992 CAPLUS
                         (4) Tackett, T; US 5117016 A 1992 CAPLUS
```

```
. . . plasma cholesterol level has a direct assocn. with coronary
heart
     disease, ischemic heart disease, acute myocardial infarction, and
     atherosclerosis. A lower cholesterol level reduces
     their risk. Among the plasma cholesterol fractions the low-d.
lipoprotein
     cholesterol and first of all its oxidized forms have strong relationship
     to the mentioned diseases. The chem. similar phytosterols in
     plants have favorable effect, they prevent the absorption of cholesterol
     from the intestine. The high-d. lipoprotein transports the cholesterol.
     . . may be influenced by nutrition. It looks advantageous the diet contg. low fat, low satd. fatty acids, low cholesterol, more
     unsatd. fatty acids, high consumption of
     fruits and vegetables, adequate intake of dietary fiber. The high-d.
     lipoprotein level will be in creased mainly.
ACCESSION NUMBER:
                         1999:521301 CAPLUS
DOCUMENT NUMBER:
                         131:228043
TITLE:
                         The essential and accursed cholesterol
AUTHOR(S):
                         Biro, Gyorqy
CORPORATE SOURCE:
                         Budapest, Hung.
                         Elelmez. Ip. (1999), 53(6), 161-166
SOURCE:
                         CODEN: EMIPAB; ISSN: 0013-5909
PUBLISHER:
                         METE
DOCUMENT TYPE:
                         Journal; General Review
LANGUAGE:
                         Hungarian
     Elelmez. Ip. (1999), 53(6), 161-166
     CODEN: EMIPAB; ISSN: 0013-5909
     ANSWER 3 OF 11 CAPLUS COPYRIGHT 2000 ACS
Ь6
ΤI
     Use of mixtures containing phytostenols for producing
     hypocholesteremic preparations
AΒ
    Mixts. of active agents contg. (a) phytostenols and/or
     phytostenol esters and (b) conjugated fatty acids or their
     glycerides are used to produce hypocholesteremic prepns. These
     mixts. have a synergistic effect in reducing the cholesterol content of
     serum. When encapsulated in gelatin, the prepns.. .
    hypocholesteremic phytostenol unsatd
     fatty acid; synergistic hypocholesteremic
     phytostenol fatty acid
     Unsaturated fatty acids
IT
     RL: BAC (Biological activity or effector, except adverse); FFD (Food or
     feed use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (diunsatd., with conjugated double bonds; use of mixts. contg.
     phytostenols for producing hypocholesteremic prepns.)
ΙT
     Sterol esters
     Sterols
     RL: BAC (Biological activity or effector, except adverse); FFD (Food or
     feed use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (from plants; use of mixts. contg. phytostenols for producing
     hypocholesteremic prepns.)
IT
     Glycerides, biological studies
     RL: BAC (Biological activity or effector, except adverse); FFD (Food or
     feed use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (polyunsatd. fatty acid-contg., with conjugated double bonds; use of
       mixts. contg. phytostenols for producing
     hypocholesteremic prepns.)
IT
    Anticholesteremic agents
     Butter
     Capsules (drug delivery systems)
     Cocoa products
     Dietary food
     Food
    Margarine
    Mayonnaise
     Salad dressings
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Sausage
     Synergistic drug interactions
        (use of mixts. contg. phytostenols for producing
      hypocholesteremic prepns.)
     Fats and Glyceridic oils, biological studies
     RL: FFD (Food or feed use); BIOL (Biological study); USES (Uses)
        (use of mixts. contg. phytostenols for producing
      hypocholesteremic prepns.)
IT
     Polyunsaturated fatty acids
     RL: BAC (Biological activity or effector, except adverse); FFD (Food or
     feed use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (with conjugated double bonds; use of mixts. contq.
     phytostenols for producing hypocholesteremic prepns.)
IT
     Fatty acid esters
     RL: BAC (Biological activity or effector, except adverse); FFD (Food or
     feed use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (with phytostenols; use of mixts. contq. phytostenols
        for producing hypocholesteremic prepns.)
ΙT
     83-45-4, .beta.-Sitostanol 83-45-4D, .beta.-Sitostanol
     , esters 83-46-5 83-46-5D, esters 1839-11-8D, 9,11-Octadecadienoic
     acid, esters with phytostenols 41005-65-6 109033-78-5
     RL: BAC (Biological activity or effector, except adverse); FFD (Food or
     feed use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (use of mixts. contg. phytostenols for producing
      hypocholesteremic prepns.)
ACCESSION NUMBER:
                         1999:344854 CAPLUS
DOCUMENT NUMBER:
                          130:347399
                         Use of mixtures containing phytostenols for
TITLE:
                         producing hypocholesteremic preparations
INVENTOR(S):
                          Fabry, Bernd
PATENT ASSIGNEE(S):
                          Henkel Kommanditgesellschaft auf Aktien, Germany
SOURCE:
                          PCT Int. Appl., 19 pp.
                          CODEN: PIXXD2
DOCUMENT TYPE:
                          Patent
LANGUAGE:
                          German
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
                                          APPLICATION NO. DATE
     PATENT NO.
                    KIND DATE
                                          WO 1998-EP7059 19981105
     WO 9925362
                      A1 19990527
         W: AU, BG, BR, BY, CA, CN, CZ, HU, ID, IS, JP, KR, LT, LV, MX, NO, NZ, PL, RO, RU, SI, SK, TR, UA, US
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
             PT, SE
     DE 19750453
                             19990527
                       A1
                                           DE 1997-19750453 19971114
                             19990607
     AU 9915603
                       A1
                                            AU 1999-15603
                                                              19981105
     EP 1028733
                       A1
                             20000823
                                            EP 1998-959848
                                                              19981105
         R: DE, ES, FR, GB, IT, NL
PRIORITY APPLN. INFO.:
                                            DE 1997-19750453 19971114
                                            WO 1998-EP7059 19981105
                         MARPAT 130:347399
OTHER SOURCE(S):
     PCT Int. Appl., 19 pp.
     CODEN: PIXXD2
REFERENCE COUNT:
REFERENCE(S):
                          (1) Funes; 1980, 5, CAPLUS
                          (2) Funes, C; AN ASOC QUIM ARGENT 1978, V66(5), P239
                          (3) Hasegawa; Hypocholesteraemic Effect of Linoleic
                              Acid and Phytosterol 1984, 25, CAPLUS
                          (4) Hasegawa; JOSHI EIYO DAIGAKU KIYO 1983, V14, P165
                              CAPLUS
                          (5) Kosbab, J; WO 9833494 A 1998
                          ALL CITATIONS AVAILABLE IN THE RE FORMAT
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```
. . onto the Indian market for human use. RBO contains oleic acid
         (38.4\%), linoleic acid (34.4\%), and linolenic acid (2.2\%) as
         unsatd. fatty acids, and palmitic (21.5%) and
         stearic (2.9\frac{1}{8}) acids as satd. fatty acids. The unsaponifiable fraction
         (4.2\%) has total tocopherols (81.3 \text{ mg}\%),. . . and 24-methylene cycloartanol (494 \text{ mg}\%). Studies on exptl. rats demonstrated a
         hypolipidemic effect of RBO. The unsaponifiable fraction of RBO
         lowers cholesterol levels. Feeding phytosterols
         , CA, and 24-methylene cycloartanol in amts. present in RBO to
         hypercholesterolemic rats for 8 wk indicates that CA alone reduces.
         and triglyceride levels significantly. Endogenous sterol excretion
         increases in animals given CA. The accumulation of CA in the liver
         inhibits cholesterol esterase activity, which in turn leads to
         redn. in circulating cholesterol levels. CA is
         structurally similar to cholesterol and may compete with the binding
    sites
         of cholesterol and sequestrate cholesterol, which. . . derivs. RBO,
         which is rich in tocopherols and tocotrienols, may improve oxidative
         stability. Tocotrienols inhibit HMG CoA reductase, resulting in
         hypocholesterolemia. The hypolipidemic effect of RBO has also
         been established in human subjects. Thus, RBO could be a suitable edible
    ACCESSION NUMBER:
                             1992:127428 CAPLUS
    DOCUMENT NUMBER:
                             116:127428
    TITLE:
                             Nutritional and biochemical aspects of the
                             hypolipidemic action of rice bran oil: a review
    AUTHOR(S):
                             Rukmini, Cheruvanky; Raghuram, Thummala C.
    CORPORATE SOURCE:
                             Natl. Inst. Nutr., Indian Counc. Med. Res.,
    Hyderabad,
                             500007, India
    SOURCE:
                             J. Am. Coll. Nutr. (1991), 10(6), 593-601
                             CODEN: JONUDL; ISSN: 0731-5724
    DOCUMENT TYPE:
                             Journal; General Review
    LANGUAGE:
                             English
         J. Am. Coll. Nutr. (1991), 10(6), 593-601
         CODEN: JONUDL; ISSN: 0731-5724
AB L6
         ANSWER 5 OF 11 MEDLINE
           . . onto the Indian market for human use. RBO contains oleic acid
         (38.4\%), linoleic acid (34.4\%), and linolenic acid (2.2\%) as
         unsaturated fatty acids, and palmitic (21.5%)
         and stearic (2.9\frac{1}{8}) acids as saturated fatty acids. The unsaponifiable
         fraction (4.2%) has total tocopherols (81.3 mg%),. . . and
    24-methylene
         cycloartanol (494 mg%). Studies on experimental rats demonstrated a
         hypolipidemic effect of RBO. The unsaponifiable fraction of RBO
         lowers cholesterol levels. Feeding phytosterols
         , CA, and 24-methylene cycloartanol in amounts present in RBO to
        hypercholesterolemic rats for 8 weeks indicates that CA alone reduces
         cholesterol and triglyceride levels significantly. Endogenous
         sterol excretion increases in animals given CA. The accumulation of CA in
         the liver inhibits cholesterol esterase activity, which in turn
         leads to reduction in circulating cholesterol levels.
         CA is structurally similar to cholesterol and may compete with
        the binding sites of cholesterol and sequestrate
        cholesterol, which is metabolized to its derivatives. RBO, which
        is rich in tocopherols and tocotrienols, may improve oxidative stability.
        Tocotrienols inhibit HMG CoA reductase, resulting in
        hypocholesterolemia. The hypolipidemic effect of RBO has also been
        established in human subjects. Thus, RBO could be a suitable edible oil.
    ACCESSION NUMBER:
                        92121598
                                     MEDLINE
   DOCUMENT NUMBER:
                        92121598
```

Nutritional and biochemical aspects of the hypolipidemic

action of rice bran oil: a review.

TITLE:

```
AUTHOR:
                     Rukmini C; Raghuram T C
                     National Institute of Nutrition, Indian Council of Medical
CORPORATE SOURCE:
                     Research, Hyderabad..
SOURCE:
                     JOURNAL OF THE AMERICAN COLLEGE OF NUTRITION, (1991 Dec)
10
                     (6) 593-601. Ref: 36
                     Journal code: H51. ISSN: 0731-5724.
PUB. COUNTRY:
                     United States
                     Journal; Article; (JOURNAL ARTICLE)
                     General Review; (REVIEW)
                     (REVIEW, TUTORIAL)
LANGUAGE:
                     English
FILE SEGMENT:
                     Priority Journals
ENTRY MONTH:
                     199204
     JOURNAL OF THE AMERICAN COLLEGE OF NUTRITION, (1991 Dec) 10 (6) 593-601.
     Journal code: H51. ISSN: 0731-5724.
L6
     ANSWER 6 OF 11 MEDLINE
     . . fraction of soybean (PUFS) given over 24 weeks to 19 patients
AΒ
     with primary types IIa and IIb hyperlipoproteinemia. The percent
     reduction of plasma cholesterol in types IIa and IIb
     were 13.2% and 11.5%, respectively. PUFS significantly reduced LDL
     cholesterol levels, but had little affect on VLDL or HDL
     cholesterol. The triglyceride/cholesterol ratio in HDL
     fraction was also significantly reduced, suggesting that PUFS plays a
     in the catabolism of HDL. PUFS contained tocopherol and
     unsaturated fatty acid. The relative impact of
     the individual components could not be assessed directly, however, it
     seemed reasonable to conclude that the hypo-cholesterolemic
     effect of PUFS results from a summation effect of plant sterols,
     tocopherols and unsaturated fatty acids.
СТ
TU, therapeutic use
      Cholesterol: BL, blood
     Hypercholesterolemia, Familial: BL, blood *Hypercholesterolemia, Familial: DT, drug therapy
      Lipoproteins, LDL Cholesterol: BL, blood
     *Phytosterols: TU, therapeutic use
      Soybeans
      Triglycerides: BL, blood
     0 (Antilipemic Agents); 0 (Lipoproteins, LDL Cholesterol); 0 (
     Phytosterols); 0 (Triglycerides)
ACCESSION NUMBER:
                    85148557
                                  MEDLINE
DOCUMENT NUMBER:
                    85148557
TITLE:
                    Effect of the purified unsaponifiable fraction of soybean
                    on primary type II hyperlipoproteinemia.
AUTHOR:
                    Nakashima Y; Nakamura T; Aramaki Y; Kuroiwa A
                    ARTERY, (1983) 12 (3) 199-211.
Journal code: 8NN. ISSN: 0098-6127.
SOURCE:
PUB. COUNTRY:
                    United States
                    Journal; Article; (JOURNAL ARTICLE)
LANGUAGE:
                    English
FILE SEGMENT:
                    Priority Journals
ENTRY MONTH:
                    198506
    ARTERY, (1983) 12 (3) 199-211.
     Journal code: 8NN. ISSN: 0098-6127.
    ANSWER 7 OF 11 EMBASE COPYRIGHT 2000 ELSEVIER SCI. B.V.
L6
     . . onto the Indian market for human use. RBO contains oleic acid
AB
     (38.4\%), linoleic aid (34.4\%), and linolenic acid (2.2\%) as
     unsaturated fatty acids, and palmitic (21.5%)
     and stearic (2.9\%) acids as saturated fatty acids. The unsaponifiable
     fraction (4.2%) has total tocopherols (81.3 mg%),. . . and
```

24-methylene

cycloartanol (494 mg%). Studies on experimental rats demonstrated a hypolipidemic effect of RBO. The unsaponifiable fraction of RBO lowers cholesterol levels. Feeding phytosterols , CA, and 24-methylene cycloartanol in amounts present in RBO to hypercholesterolemic rats for 8 weeks indicates that CA alone reduces cholesterol and triglyceride levels significantly. Endogenous sterol excretion increases in animals given CA. The accumulation of CA in the liver inhibits cholesterol esterase activity, which in turn leads to reduction in circulating cholesterol levels. CA is structurally similar to cholesterol and may compete with the binding sites of cholesterol and sequestrate cholesterol, which is metabolized to its derivatives. RBO, which is rich in tocopherols and tocotrienols, may improve oxidative stability. Tocotrienols inhibit HMG CoA reductase, resulting in hypocholesterolemia. The hypolipidemic effect of RBO has also been established in human subjects. Thus, RBO could be a suitable edible oil.

ACCESSION NUMBER: 92262702 EMBASE

DOCUMENT NUMBER: 1992262702

TITLE: Nutritional and biochemical aspects of the hypolipidemic

action of rice bran oil: A review.

AUTHOR: Rukmini C.; Raghuram T.C.

CORPORATE SOURCE: National Institute of Nutrition, Hyderabad 500007, India

SOURCE: Journal of the American College of Nutrition, (1991) 10/6

(593-601).

ISSN: 0731-5724 CODEN: JONUDL

COUNTRY: United States

DOCUMENT TYPE: Journal; General Review FILE SEGMENT: 006 Internal Medicine

Public Health, Social Medicine and Epidemiology
Cardiovascular Diseases and Cardiovascular Surgery

029 Clinical Biochemistry

LANGUAGE: English SUMMARY LANGUAGE: English

SO Journal of the American College of Nutrition, (1991) 10/6 (593-601). ISSN: 0731-5724 CODEN: JONUDL

L6 ANSWER 8 OF 11 WPIDS COPYRIGHT 2000 DERWENT INFORMATION LTD

TI Oil and fat composition containing **phytosterol** for preparation of food and pharmaceuticals.

AB WO 9959423 UPAB: 20000330

NOVELTY - Composition comprises a **phytosterol** dissolved in an oil and fat comprising one or more polyhydric alcohol/fatty acid esters each having an esterification degree of. . . more and containing at least one unesterified hydroxyl group.

DETAILED DESCRIPTION - An INDEPENDENT CLAIM is included for a **phytosterol** containing food which comprises one or more lipid ingredients comprising at least 15 wt.% one or more diacylglycerols. The weight ratio of diacylglycerols to **phytosterol** is 10-100.

USE - Used for preparation of oil in water type fat-processed foods such as drinks, desserts, ice creams,. . . cakes, cookies, pies, breads

and chocolates, and other foods including bakery mixes, processed meat products, frozen entrees and frozen foods, **hypocholesteremic** preparation in the form of a capsule, sugar-coated tablet, molded granules, candy or drop.

ADVANTAGE - Blood cholesterol level is. . . good workability, and the flavor and feeling of food items are not changed. Generation of smoking during cooking is eliminated. **Phytosterol** is solubilized by the composition.

Dwg.0/0

TECH.

esterification degree of 2-4.5 and comprises glycerol/fatty acid esters, polyglycerol/fatty acid esters, sucrose/fatty acid esters or sorbitan/fatty acid esters. The **phytosterol** is contained in an

amount of 1.2 wt.% or more. 55 wt.% Fatty acids contained in diacylglycerols are unsaturated fatty acids.

The composition is a frying oil.

TT: OIL FAT COMPOSITION CONTAIN **PHYTOSTEROL** PREPARATION FOOD. PHARMACEUTICAL.

ACCESSION NUMBER: 2000-105555 [09] WPIDS

CROSS REFERENCE: 1999-571935 [48] DOC. NO. CPI: C2000-031612

TITLE: Oil and fat composition containing phytosterol

for preparation of food and pharmaceuticals.

DERWENT CLASS: A96 B01 D13

INVENTOR(S): GOTO, N; NISHIDE, T; TANAKA, Y; YASUKAWA, T

PATENT ASSIGNEE(S): (KAOS) KAO CORP

COUNTRY COUNT: 23

PATENT INFORMATION:

PA	ГЕИТ	ИО	KIND	DATE	WEEK	LA	PG
WO	9959	9423	A1	19991125	(200009)*	EN	26

RW: AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE

W: BR CA CN SG

US 6025348 A 20000215 (200016)#

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 9959423	A1	WO 1998-JP2228	19980521
US 6025348	А	US 1998-69754	19980430

PRIORITY APPLN. INFO: WO 1998-JP2228 19980521; US 1998-69754 19980430

- L6 ANSWER 9 OF 11 WPIDS COPYRIGHT 2000 DERWENT INFORMATION LTD

 TI . . . risk of cardiovascular or lipid diseases, diabetes and thyroid disease using as risk indicator the ratio of campesterol to betasitosterol in the serum, also treatment of these disorders with phytosterol(s).
- AB WO 9801759 UPAB: 19980302

 The ratio (R), in serum, of campesterol (I) to beta -sitosterol

 (II) is determined, compared with that in a normal control and used as an indicator of increased risk of cardiovascular. . . Also claimed are:

 (1) method for assessing risk of cardiovascular and lipid disorders by measuring (R), serum levels of total phytosterols (TP) and total cholesterol (TC) and comparing all 3 values with those of normal controls;
- (2) method for increasing the inhibitory effect of phytosterols (III) on enterocyte absorption of cholesterol (IV) by administering (III) that inhibits absorption of (IV) and/or (II); (3) (IV)-lowering composition. . . treatment of diabetes and thyroid disease or dietary modification to increase TP and (R). (III) are hydrophobic, particularly (I) and sitosterol, and may be supplied as safflower, sesame seed, maize, rice bran, olive, rapeseed, flaxseed or coconut oils. Particularly (III) are administered together with (i) at least 1 inhibitor
- of (IV) biosynthesis, especially (A); (ii) one or more saturated or mono/poly-unsaturated fatty acids (also providing a synergistic effect) or (iii) plant oils as specified above. Optionally the composition of (3) also includes 2-6%. . . of the (IV) level, e.g. in subjects with apparently normal (IV) level. (III) lowers serum TC and low-density lipoprotein (LDL) cholesterol, but increases HDL and the HDL/LDL ratio, and there is a synergistic effect when used with inhibitors (A) of 3-hydroxy-3-methylglutaryl coenzyme A reductase, allowing a reduction in the dose of (A), and

thus of side effects.

Dwg.1/13

TT: DIAGNOSE RISK CARDIOVASCULAR LIPID DISEASE DIABETES THYROID DISEASE

RISK INDICATE RATIO BETA SITOSTEROL SERUM TREAT DISORDER

PHYTOSTEROL.

ACCESSION NUMBER:

1998-101200 [09] WPIDS

DOC. NO. NON-CPI: DOC. NO. CPI:

N1998-081048 C1998-033490

TITLE:

Diagnosing risk of cardiovascular or lipid diseases, diabetes and thyroid disease - using as risk indicator

the ratio of campesterol to beta-sitosterol in the serum, also treatment of these disorders with

phytosterol(s).

DERWENT CLASS:

B04 D16 S03

INVENTOR(S):

NOVAK, E

PATENT ASSIGNEE(S):

(FORB-N) FORBES MEDI-TECH INC

COUNTRY COUNT:

PATENT INFORMATION:

PATENT NO	KIND DATE	WEEK	LA	PG

WO 9801759 A1 19980115 (199809) * EN 93

RW: AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE

W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GE GH HU IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK TJ TM TR TT UA UG UZ VN YU

AU 9732518 A 19980202 (199826)

EP 912900 A1 19990506 (199922)

R: AL AT BE CH DE DK ES FI FR GB GR IE IT LI LT LU LV MC NL PT RO SE ST

US 5965449 A 19991012 (199949)

CZ 9900011 A3 20000112 (200009)

HU 9904042 A2 20000328 (200025)

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 9801759	A1	WO 1997-CA474	19970703
AU 9732518	A	AU 1997-32518	19970703
EP 912900	A1	EP 1997-928091	19970703
		WO 1997-CA474	19970703
US 5965449	A	US 1996-675018	19960703
CZ 9900011	A3	WO 1997-CA474	19970703
		CZ 1999-11	19970703
HU 9904042	A2	WO 1997-CA474	19970703
		HU 1999-4042	19970703

FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 9732518	A Based on	WO 9801759
EP 912900	Al Based on	WO 9801759
CZ 9900011	A3 Based on	WO 9801759
HU 9904042	A2 Based on	WO 9801759

PRIORITY APPLN. INFO: US 1996-675018 19960703

H 16 ANSWER 10 OF 11 BIOSIS COPYRIGHT 2000 BIOSIS

. . onto the Indian market for human use. RBO contains oleic acid (38.4%), linoleic acid (34.4%), and linolenic acid (2.2%) as unsaturated fatty acids, and palmitic (21.5) and stearic (2.9%) acids as saturated fatty acids. The unsaponifiable fraction (4.2%) has total tocopherols (81.3% mg%),. . and

of

RBO lowers cholesterol levels. Feeding phytosterols, CA, and 24-methylene cycloartanol in amounts present in RBO to hypercholesterolemic rats for 8 weeks indicates that CA alone reduces cholesterol and triglyceride levels significantly. Endogenous sterol excretion increases in animals given CA. The accumulation of CA in the liver inhibits cholesterol esterase activity, which in turn leads to reduction in circulating cholesterol levels. CA is structurally similar to cholesterol and may compete with the binding sites of cholesterol and sequestrate cholesterol, which is metabolized to its derivatives. RBO, which is rich in tocopherols and tocotrienols, may improve oxidative stability. Tocotrienols inhibit HMG COA reductase, resulting in hypocholesterolemia. The hypolipidemic effect of RBO has also been established in human subjects. Thus, RBO could be a suitable edible oil. . .

ACCESSION NUMBER: 1992:75496 BIOSIS

DOCUMENT NUMBER:

BA93:43951

TITLE:

NUTRITIONAL AND BIOCHEMICAL ASPECTS OF THE HYPOLIPIDEMIC

ACTION OF RICE BRAN OIL A REVIEW.

AUTHOR(S):

RUKMINI C; RAGHURAM T C

CORPORATE SOURCE:

NATL. INST. NUTRITION, HYDERABAD 500007, INDIA.

SOURCE:

J AM COLL NUTR, (1991) 10 (6), 593-601. CODEN: JONUDL. ISSN: 0731-5724.

FILE SEGMENT:

BA; OLD

LANGUAGE:

English

SO J AM COLL NUTR, (1991) 10 (6), 593-601.

CODEN: JONUDL. ISSN: 0731-5724.

L6 ANSWER 11 OF 11 BIOSIS COPYRIGHT 2000 BIOSIS

AB The effects of dietary fats and **phytosterol** on the fatty acid composition and lipoprotein cholesterol in serum were studied in female rats with the following results. The. . . due to an increase in the lower density lipoprotein (LDL + VLDL [very low density lipoprotein]).

The

addition of 5% **phytosterol** to the 10% butter-cholesterol diet reduced the total cholesterol level and increased the ratio of cholesterol

in high density lipoprotein. . . = 0.947), and also the level of LDL + VLDL-cholesterol (r = 0.935). Cod liver oil, wheat germ oil and phytosterol induce an increase in the PUFA/SFA ratio, promote hypocholesterolemia and change lipoprotein concentration. There were indications that no relationship exists between the change in the total cholesterol level and . . .

IT Miscellaneous Descriptors

COD LIVER OIL WHEAT GERM OIL POLY UNSATURATED FATTY

-ACID SATURATED FATTY-ACID HYPO CHOLESTEROLEMIA OLEIC-ACID

ACCESSION NUMBER:

1984:349467 BIOSIS

DOCUMENT NUMBER:

BA78:85947

TITLE:

EFFECTS OF DIETARY FATS AND PHYTO STEROL ON SERUM

FATTY-ACID COMPOSITION AND LIPO PROTEIN CHOLESTEROL IN

RATS.

AUTHOR(S):

HIRAI K; OHNO Y; NAKANO T; IZUTANI K

CORPORATE SOURCE:

DEP. NUTR. BIOCHEM., FAC. SCI. LIVING, OSAKA CITY UNIV.,

SUMIYOSHI, OSAKA 558, JPN.

SOURCE:

J NUTR SCI VITAMINOL, (1984) 30 (2), 101-112.

CODEN: JNSVA5. ISSN: 0301-4800.

FILE SEGMENT:

BA; OLD

LANGUAGE:

English

SO J NUTR SCI VITAMINOL, (1984) 30 (2), 101-112.